

## SEQUENCE LISTING

<110> TRANSGENE S.A.

<120> Poxvirus with targeted infection specificity

<130> D18836

<150> EP 00 44 0109

<151> 2000-04-14

<150> EP 01 44 0009

<151> 2001-01-22

<150> US 60/246 080

<151> 2000-11-07

<160> 21

<170> PatentIn Ver. 2.1

<210> 1

<211> 24

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR primer to amplify the MVA 138L gene and flanking region

<400> 1

cagactggac ggcgtccata tgag

24

<210> 2

<211> 61

<212> DNA

<213> Artificial Sequence

<220>

<221> gene

<222> Complement((1)..(61))

<220>

<223> Description of Artificial Sequence: antisens PCR primer to amplify the 3' end of MVA 138L gene and 3' flanking region

<400> 2

catTTTTTaa gtatagaata aaagatcccg ggagtaccat cgtgattctt accagatatt 60  
a 61

<210> 3

<211> 61

<212> DNA

<213> Artificial Sequence

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<220>
<223> Description of Artificial Sequence: PCR primer to
      amplify E. coli gpt gene and H5R promoter

<220>
<221> gene
<222> (1)..(61)

<400> 3
taatatctgg taagaatcac gatggtactc cgggcatctt ttattctata cttaaaaaat 60
g                                                    61

<210> 4
<211> 35
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence: antisense PCR
      primer to amplify E. coli GPT gene and pH5R
      promoter

<400> 4
gggggtaatt aaggaagtta aaaagaacaa cgccc                                35

<210> 5
<211> 38
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence: PCR primer to
      amplify the upstream region of MVA 138L gene.

<400> 5
gggggaattc gagcttatag cgtttagttc aggtacgg                                38

<210> 6
<211> 44
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence: antisense PCR
      primer to amplify the upstream region of the MVA
      138L gene

<400> 6
ggggaagctt ttaaagtaca gattttagaa actgacactc tgcg                                44

<210> 7
<211> 68
<212> DNA

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<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: antisense  
primer to amplify the upstream region of the MVA  
138L gene

<400> 7

ggggaagctt caagagcggc acggctcccg ccgctgcgac gttcaggagg accaaggcaa 60  
ccacgaac 68

<210> 8

<211> 31

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR primer to  
amplify the MVA 138L gene and its downstream  
region

<400> 8

ggggaagctt atggacggaa ctcttttccc c 31

<210> 9

<211> 37

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: antisense PCR  
primer to amplify the MVA 138L gene and its  
downstream region

<400> 9

gggggaattc gcttatcggt atcggttcta gcttctg 37

<210> 10

<211> 68

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR primer to  
amplify SM3 scFv sequence

<400> 10

cgcagagtgt cagtttctaa aatctgtact ttaaatgggt cagctgcagg agtctggagg 60  
aggcttgg 68

<210> 11

<211> 58

<212> DNA

<213> Artificial Sequence

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<220>
<223> Description of Artificial Sequence: antisense PCR
        primer to amplify the SM3 scFv sequence

<400> 11
gatcgatcatc tccggggaaa agagttccgt ccatacagttt ggttcctcca ccgaacac   58

<210> 12
<211> 57
<212> DNA
<213> Artificial Sequence

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<223> Description of Artificial Sequence: PCR primer to
        amplify the SM3 scFv sequence

<400> 12
cctgaacgtc gcagcggcgg gagccgtgcc gctcttggtg cagctgcagg agtctgg   57

<210> 13
<211> 111
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence: sequence of
        the synthetic plik7.5 promoter

<400> 13
ataaaaaatat agtagaatTT cATTtGTTTT tttctatgct ataaatagga tccgataaag 60
tgaaaaataa ttctaattta tgcacggta aggaagtaga atcataaaga a          111

<210> 14
<211> 53
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence:PCR primer to
        amplify the plik7.5 promoter

<400> 14
gggggatccc ccgggctgca gaagcttttc ttatgattc tacttctta ccg          53

<210> 15
<211> 50
<212> DNA
<213> Artificial Sequence

<220>
<223> Description of Artificial Sequence: antisense PCR
        primer to amplify the plik7.5 promoter

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<400> 15  
 ggggggagat ctaagcttgt cgacataaaa atatagtaga atttcatttg 50

<210> 16  
 <211> 77  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: synthetic  
 sequence

<400> 16  
 gatggtgaca gggggaatgg caagcaagtg ggatctcgag ttgggtgact ttggtgacag 60  
 atactactgt gtttaag 77

<210> 17  
 <211> 85  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: synthetic  
 sequence

<400> 17  
 gatccttaaa cacagtagta tctgtcacca aagtcaccca actcgagatc ccacttgctt 60  
 gccattcccc ctgtcccat ctgca 85

<210> 18  
 <211> 32  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:PCR primer to  
 amplify the 5' F13L flanking region of MVA

<400> 18  
 gagaggatcc gggatatctag ccacagtaaa tc 32

<210> 19  
 <211> 32  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:Description of  
 Artificial Sequence :antisense PCR primer to  
 amplify the 5' F13L flanking region of MVA

<400> 19  
 ttctgaattc ggaatctgta ttctcaatc cg 32

<210> 20  
 <211> 33  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: PCR primer to  
 amplify the 3' F13L flanking region of MVA

<400> 20  
 atctgaattc gtggagatga tgatagttta agc

33

<210> 21  
 <211> 34  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: antisense PCR  
 primer to amplify the 3' F13L flanking region of  
 MVA

<400> 21  
 aacaggatcc cttatacatc ctgttctatc aacg

34